PATENT COOPERATION TREATY

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INTERNATIONAL SEARCHING AUTHORITY To: **BORDEN LADNER GERVAIS LLP** World Exchange Plaza WRITTEN OPINION OF THE 1100 - 100 Oueen Street INTERNATIONAL SEARCHING AUTHORITY OTTAWA, Ontario Canada, K1P 1J9 (PCT Rule 43bis.1) Date of mailing 06 June 2005 (06.06.2005) (day/month/year) Applicant's or agent's file reference FOR FURTHER ACTION PAT2786W-90 See paragraph 2 below International application No. International filing date (day/month/year) Priority date (day/month/year) 02 March 2004 (02-03-2004) 02 March 2005 (02-03-2005) PCT/CA2005/000323 International Patent Classification (IPC) or both national classification and IPC A61K 31/167 A61P 29/00 A61P 11/00 A61P 31/00 **Applicant** MCGILL UNIVERSITY ET AL 1. This opinion contains indications relating to the following items: [X] Box No. I Basis of the opinion [X] Box No. II **Priority** [X] Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability [] Box No. IV Lack of unity of invention [X] Box No. V Reasoned statement under Rule 43bis. 1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement. Box No. VI Certain documents cited Box No. VII Certain defects in the international application [X] Box No. VIII Certain observations on the international application 2. FURTHER ACTION If a demand for international preliminary examination is made, this opinion will be considered to be a written opinion of the International Preliminary Examining Authority ("IPEA") except that this does not apply where the applicant chooses an Authority other than this one to be the IPEA and the chosen IPEA has notified the International Bureau under Rule 66.1bis(b) that written opinions of this International Searching Authority will not be so considered. If this opinion is, as provided above, considered to be a written opinion of the IPEA, the applicant is invited to submit to the IPEA a written reply together, where appropriate, with amendments, before the expiration of 3 months from the date of mailing of Form PCT/ISA/220 or before the expiration of 22 months from the priority date, whichever expires later. For further options, see Form PCT/ISA/220. 3. For further details, see notes to Form PCT/ISA/220. Authorized officer Name and mailing address of the ISA/CA Canadian Intellectual Property Office Place du Portage I, C114 - 1st Floor, Box PCT Tania Nish (819) 934-3592

50 Victoria Street

Gatineau, Quebec K1A 0C9
Facsimile No.: 001(819)953-2476

International application No. PCT/CA2005/000323

| 1. With regard to the language, this opinion has been established on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item. [] This opinion has been established on the basis of a translation from the original language into the following language which is the language of a translation furnished for the purposes of international search (under Rules 12.3 and 23.1(b)). 2. With regard to any nucleotide and/or amileo acid sequence disclosed in the international application and necessary to the claimed invention, this opinion has been established on the basis of: a. type of material [] a sequence listing [] a sequence listing [] in owritten format [] in contained in the international application as filed. [] filed together with the international application in computer readable form. [] filed together with the international application in computer readable form. [] filed together with the international application in computer readable form. [] In addition, in the case that more than one version or copy of a sequence listing and/or table relating thereto has been filed or furnished, the required statement that the information in the subsequent or additional copies is identical to that in the application as filed, as appropriate, were furnished. 4. Additional comments: | Bo | x N | io. | I Basis of this opinion | | | |
|--|----|-----|------|---|-----|--|--|
| which is the language of a translation furnished for the purposes of international search (under Rules 12.3 and 23.1(b)). 2. With regard to any nucleotide and/or amino acid sequence disclosed in the international application and necessary to the claimed invention, this opinion has been established on the basis of: a. type of material [] a sequence listing [] table(s) related to the sequence listing b. format of material [] in written format [] in computer readable form c. time of filing/furnishing [] contained in the international application as filed. [] filed together with the international application in computer readable form. [] furnished subsequently to this Authority for the purposes of search. 3 [] In addition, in the case that more than one version or copy of a sequence listing and/or table relating thereto has been filed or furnished, the required statement that the information in the subsequent or additional copies is identical to that in the application as filed or does not go beyond the application as filed, as appropriate, were furnished. | 1. | | | | | | |
| (under Rules 12.3 and 23.1(b)). 2. With regard to any nucleotide and/or amino acid sequence disclosed in the international application and necessary to the claimed invention, this opinion has been established on the basis of: a. type of material [] a sequence listing [] table(s) related to the sequence listing b. format of material [] in written format [] in computer readable form c. time of filing/furnishing [] contained in the international application as filed. [] filed together with the international application in computer readable form. [] furnished subsequently to this Authority for the purposes of search. 3 [] In addition, in the case that more than one version or copy of a sequence listing and/or table relating thereto has been filed or furnished, the required statement that the information in the subsequent or additional copies is identical to that in the application as filed, as appropriate, were furnished. | | [|] | This opinion has been established on the basis of a translation from the original language into the following language | | | |
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| b. format of material [] in written format [] in computer readable form c. time of filing/furnishing [] contained in the international application as filed. [] filed together with the international application in computer readable form. [] furnished subsequently to this Authority for the purposes of search. 3 [] In addition, in the case that more than one version or copy of a sequence listing and/or table relating thereto has been filed or furnished, the required statement that the information in the subsequent or additional copies is identical to that in the application as filed or does not go beyond the application as filed, as appropriate, were furnished. | | | | [] a sequence listing | | | |
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| filed or does not go beyond the application as filed, as appropriate, were furnished. | 3 | [|] | In addition, in the case that more than one version or copy of a sequence listing and/or table relating thereto has been filed or | | | |
| 4. Additional comments: | • | | | | 8.5 | | |
| | 4. | Ad | diti | ional comments: | | | |
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International application No. PCT/CA2005/000323

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|-----------------|---------------------------|---|--|--|--|----------------------|
| Box N | o. II | Priority | | | | |
| | | | | | | |
| 1. [7 | The follow | wing document has not yet be | en furnished: | | | |
| | [X] cop | y of the earlier application | whose priority has be | en claimed (Rules 43bis.) | and 66.7(a)). | |
| | [] tran | slation of the earlier applicat | ion whose priority has be | en claimed (Rules 43 <i>bis.</i> 1 a | nd 66.7(b)) . | |
| | | ently it has not been possi ed on the assumption that | | | This opinion has neverthe | eless been |
| 2. [| found in | nion has been established a valid (Rules 43 <i>bis</i> .1 and 6 considered to be the relev | 4.1). Thus for the purp | | | |
| 3. Ad | litional obser | rvations, if necessary: | | | | |
| es fil th | ablishing thing date of t | ocument pertaining to the nis first written opinion. He the priority document. If it nal search report could bec | ence, it is based on the t later turns out that thi | assumption that all clain is incorrect, the docume | ns enjoy priority rights fro ant, WO 2004/064823 A1 | om the , cited in |
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International application No. PCT/CA2005/000323

| Box No. | III Non-establishment | of opinion with regard to novelty, inventive step and industrial applicability |
|---------|--|---|
| | tions whether the claimed inve e have not been examined in re | ntion appears to be novel, to involve an inventive step (to be non obvious), or to be industrially spect of: |
| [] | the entire international applic | ation . |
| [X] | claim Nos. 1-5 | |
| beca | us e : | |
| ſΧΊ | the said international applicat | ion, or the said claim Nos. 1-5 |
| [] | | matter which does not require an international preliminary examination (specify): |
| | Although claims 1-3 are direct on the alleged effects of fenre | edical treatment. (Rule 39.1(iv) PCT) ted to methods of medical treatment of the human/animal body, the search has been carried out based tinide and derivatives/analogs and ceramide and derivatives/analogs on pro-inflammation, oliferation and respiratory tract infection |
| | | -5 is directed to a method of medical treatment of the human or animal body (Rule 39.1(iv)PCT). No T Contracting States for the assessment of the industrial applicability of claims 1-5 (Article |
| [X] | the description, claims or drav | vings (Indicate particular elements below) or said claim Nos. 1-5 |
| | • | gful opinion could be formed (specify): |
| | cover an extremely large number of options of comeaningful search is not possi | method to treat inflammation with a compound that increases ceramide levels in the cell. The claims ber of possible compounds, including currently undiscovered compounds. As such, the extremely impounds renders the claims unclear and not concise within the meaning of Article 6 PCT that a lible. Consequently, the search has been carried out for only a limited number of compounds which are mely fenretinide and derivatives/analogs thereof and ceramide and derivatives/analogs thereof. |
| | | |
| [] | the claims, or said claims Nos | . are so inadequately supported uningful opinion could be formed. |
| | | |
| | • | has been established for said claims Nos. |
| l J | | cid sequence listing does not comply with the standard provided for in Annex C of the |
| | Administrative Instructions in | |
| | the written form | [] has not been furnished |
| | | [] does not comply with the standard |
| | the computer readable form | [] has not been furnished |
| | | [] does not comply with the standard |
| [] | the tables related to the nucleo | otide and/or amino acid sequence listing, if in computer readable form only, do not comply with the |
| | ed for in Annex C-bis of the Administrative Instructions. | |
| | | |
| [] | See Supplemental Box for fun | her details. |

IAP9 Rec'd PCT/PTO 3 1 AUG 2006

WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No. PCT/CA2005/000323

| Box No. V | Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industria | | |
|-----------|---|--|--|
| | applicability; citations and explanations supporting such statement | | |

1. S

| Sta | tement | | | | |
|-----|-------------------------------|--------|------------|-----|--|
| | Novelty (N) | Claims | | YES | |
| | | Claims | <u>1-5</u> | ИО | |
| | Inventive step (IS) | Claims | | YES | |
| | | Claims | <u>1-5</u> | ИО | |
| | Industrial applicability (IA) | Claims | | YES | |
| | | Claims | | МО | |
| | | | | | |

2. Citations and explanations:

D1: CA 2 365 290 A1 D2: WO 01/72701 A1

D3: Am. J. Respir. Cell Mol. Biol., 22, 460-468

D4: WO 00/00207 A1

D5: J. Biol. Chem., 277, 49531-49537

D6: JNCI, 91, 1138-1146 D7: WO 2004/0644823 A1

NOVELTY

1) D1 discloses the use of ceramides, derivatives and/or precursors of ceramides in the treatment of cystic fibrosis and associated diseases and illnesses, such as inflammation and respiratory infection.

D2 discloses the use of ceramide and derivatives thereof in the prevention of cellular proliferation, inflammatory disease or inflammation.

Therefore, since the subject matter of claims 1-2 and 4-5 is the same as the subject matter disclosed by D1 and D2, claims 1-2 and 4-5 would not be considered novel with respect to D1 and D2. (Art.33(2), PCT)

2) D3 discloses the use of ceramides and analogs thereof in the mediation of cell death by apoptosis. Ceramide is disclosed as a second messanger in initiating the apoptotic response.

D4-D6 disclose the use of fenretinide and other such retinoic acid derivatives in the treatment of hyperproliferative disorders by manipulation of the ceramide-mediated apoptosis.

Therefore, since the subject matter of claim 3 is the same as the subject matter disclosed by D3-D6, claim 3 would not be considered novel with respect to D3-D6 (Art.33(2), PCT)

INVENTIVE STEP

One skilled in ther art with regards to D1-D6 would be able to conclude that the increase of ceramide levels will inhibit inflammation and proliferation and thus use of a medicament that increases ceramide levels, such as fenretinide and other like retinoic acid derivatives and precursors/derivatives of ceramide, will treat inflammation and proliferation. (Art. 33(3), PCT)

International application No. PCT/CA2005/000323

Box No. VIII Certain observations on the international application

The following observations on the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

Claims 1-5 do lack clarity and conciseness and do not comply with Article 6 (PCT) for the following reasons:

- 1) Claims 1-5 defines the compound used in the treatment as "an agent that increases ceramide levels in the cell". The definition attempts to define the therapeutic compound of the invention solely by the result to be achieved, i.e. the increase of ceramide levels in the cell. Therefore rendering the claims unclear, as it directs to a desired results rather than to the combination necessary to achieve the result as described in the description.
- 2) Claims 1-2 and 4-5 define the use of an agent to treat inflammatory response, proliferation and reduction of respiratory tract infections. The description specifically supports the treatment of respiratory inflammation, proliferation and respiratory infection in subjects with cystic fibrosis (CF) disease using a specific agent, namely fearetinide. Therefore, the claims are broader than the scope of the invention disclosed in the description.
- 3) The term "diseased cell" in claims 1 and 4 cause the claims to be ambiguous
- 4) Claim 3 defines the use of an agent to induce an inflammatory response in a cell. The claim is broader than the scope of the invention as taught in the description where fenretidine is shown to induce an inflammatory response in cells that are not in a pro-inflammatory response, ie cell which are not stimulated.